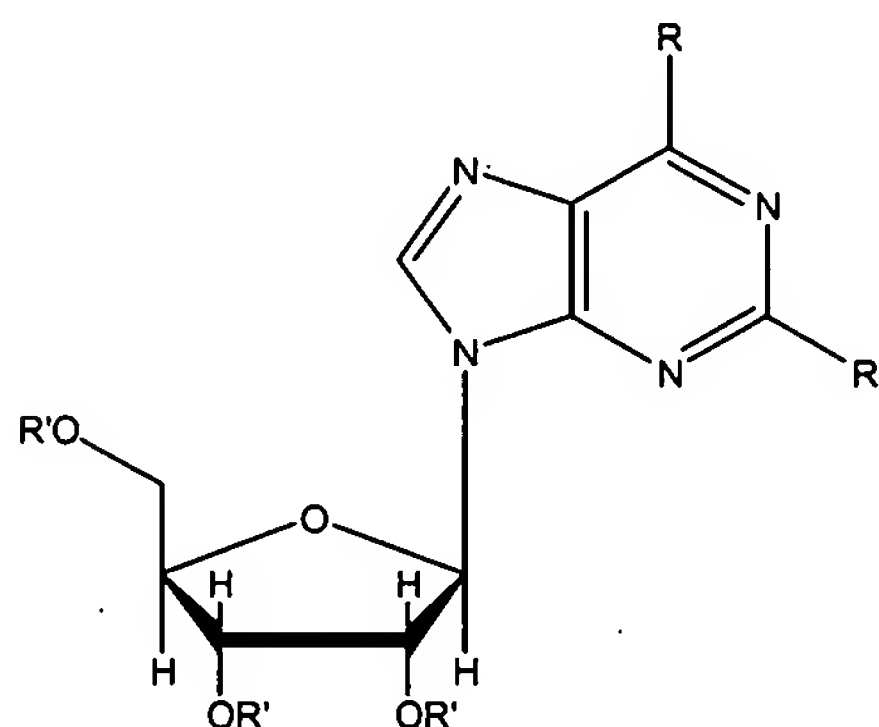


Amendments to the Claims:

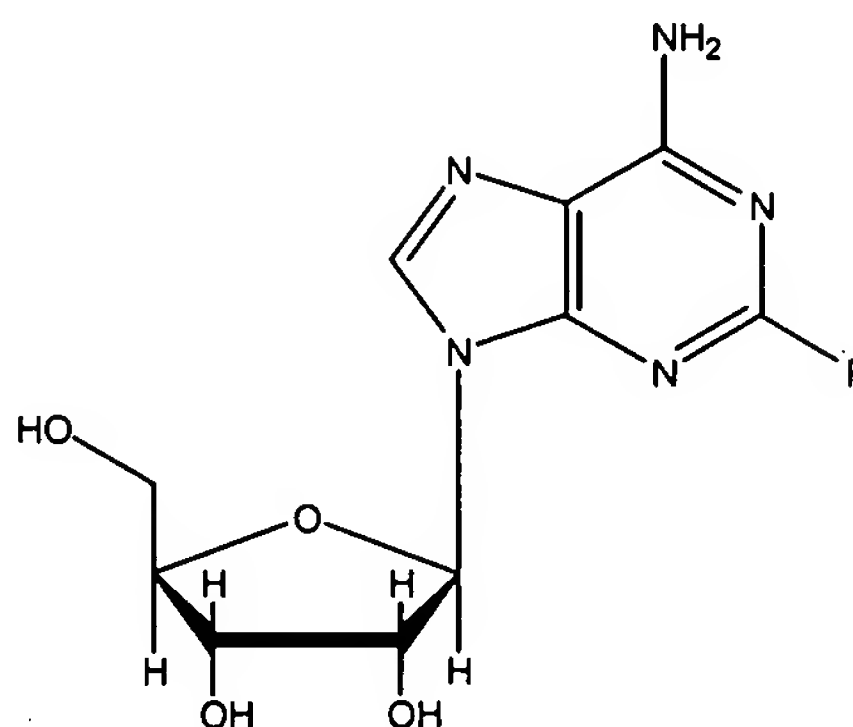
This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

1. (Original) A method of synthesis of a 2-substituted adenosine of formula I which comprises converting a compound of formula II to a compound of formula I:



II



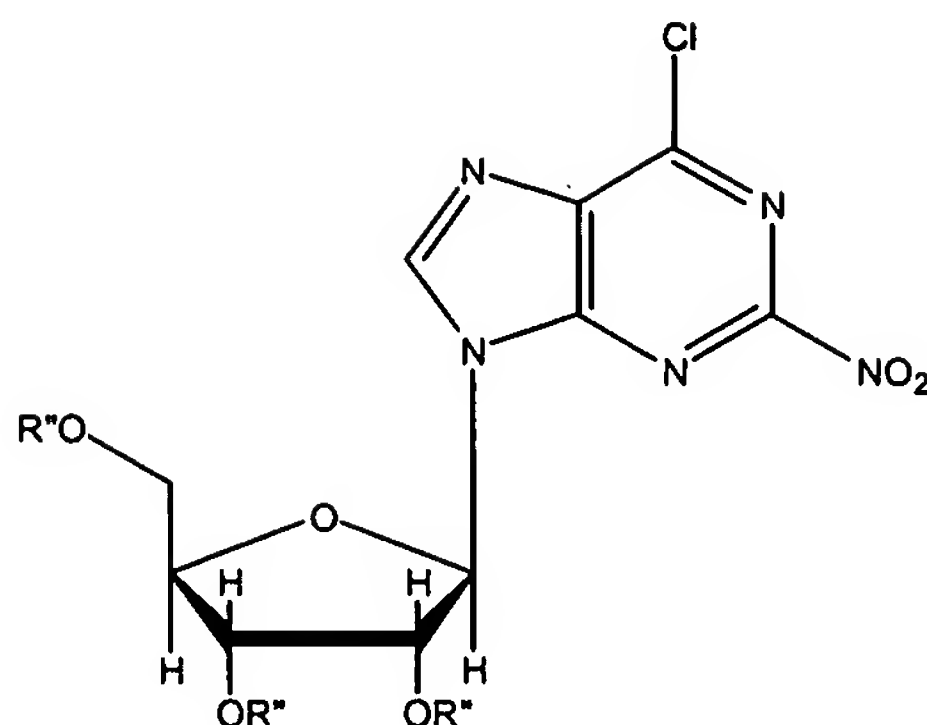
I

wherein:

R is C<sub>1-6</sub> alkoxy (straight or branched), a phenoxy group (unsubstituted, or mono-, or di-substituted by halo, amino, CF<sub>3</sub>-, cyano, nitro, C<sub>1-6</sub> alkyl, or C<sub>1-6</sub> alkoxy), a benzyloxy group (unsubstituted, or mono-, or di-substituted by halo, amino, CF<sub>3</sub>-, cyano, nitro, C<sub>1-6</sub> alkyl, or C<sub>1-6</sub> alkoxy), or a benzoyl group (unsubstituted, or mono-, or di-substituted by halo, amino, CF<sub>3</sub>-, cyano, nitro, C<sub>1-6</sub> alkyl, or C<sub>1-6</sub> alkoxy);

R' = H, or a protecting group.

2. (Original) A method according to claim 1, wherein R = methoxy, ethoxy, propoxy, butoxy, pentyloxy, hexyloxy, phenoxy, benzyloxy, or benzoyl.
3. (Currently Amended) A method according to claim 1 ~~or 2~~, wherein R' is a protecting group that can be removed under conditions that replace the R group with an amino group at the 6-position of the purine component of the compound of formula II.
4. (Original) A method according to claim 3, wherein the compound of formula II is converted to the compound of formula I in a single reaction step.
5. (Currently Amended) A method according to claim 1 ~~any preceding claim~~, wherein the protecting group is acetyl or benzoyl, and the compound of formula II is converted to the compound of formula I by treatment with ammonia.
6. (Currently Amended) A method according to claim 1 ~~or 2~~, wherein R' is H, and the compound of formula II is aminated to form the compound of formula I.
7. (Original) A method according to claim 6, wherein the compound of formula II is aminated by heating the compound in a solution of ammonia and then cooling the solution to precipitate the compound of formula I.
8. (Currently Amended) A method according to claim 1 ~~any preceding claim~~, which further comprises isolating the compound of formula I produced.
9. (Currently Amended) A method according to claim 1 ~~any preceding claim~~, which further comprises converting a compound of formula III to a compound of formula II:



**III**

wherein R'' is a protecting group, preferably acetyl or benzoyl.

10. (Original) A method of synthesis of a compound of formula II which comprises converting a compound of formula III to the compound of formula II.

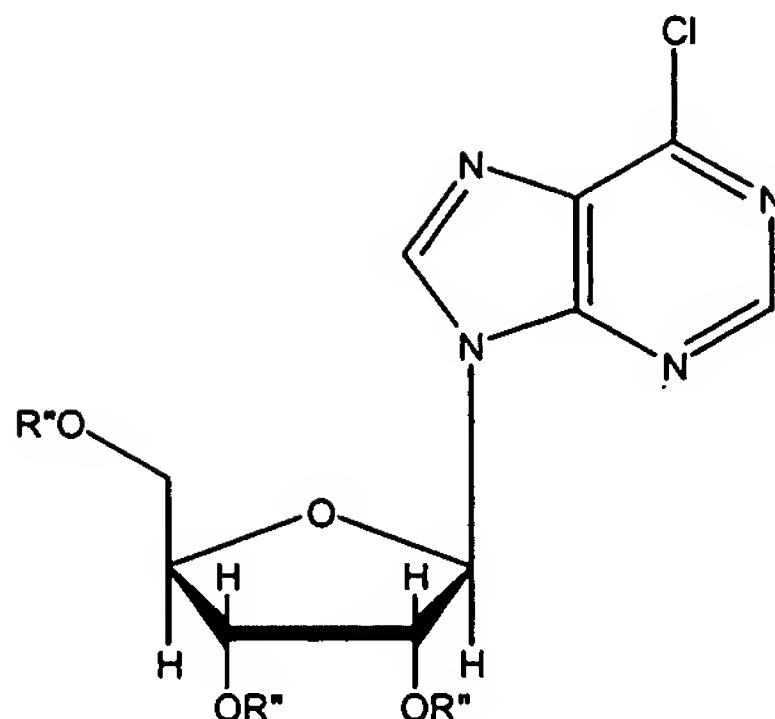
11. (Currently Amended) A method according to claim 9 ~~or 10~~, wherein the compound of formula III is alkoxylated or benzoylated to form the compound of formula II.

12. (Currently Amended) A method according to claim 9 ~~any of claims 9 to 11~~, wherein the compound of formula III is triacetoxo 2-nitro-6-chloroadenosine.

13. (Original) A method according to claim 12, wherein triacetoxo 2-nitro-6-chloroadenosine is methoxylated using sodium methoxide in methanol as methoxylating reagent.

14. (Currently Amended) A method according to claim 9 ~~any of claims 9 to 13~~, which further comprises isolating the compound of formula II produced.

15. (Currently Amended) A method according to claim 9 ~~any of claims 9 to 14~~, which further comprises converting a compound of formula IV to a compound of formula III:



IV

wherein R'' is a protecting group, preferably acetyl or benzoyl.

16. (Original) A method according to claim 15, wherein the compound of formula IV is nitrated to form the compound of formula III.

17. (Currently Amended) A method according to claim 15 ~~or 16~~, which further comprises isolating the compound of formula III produced.

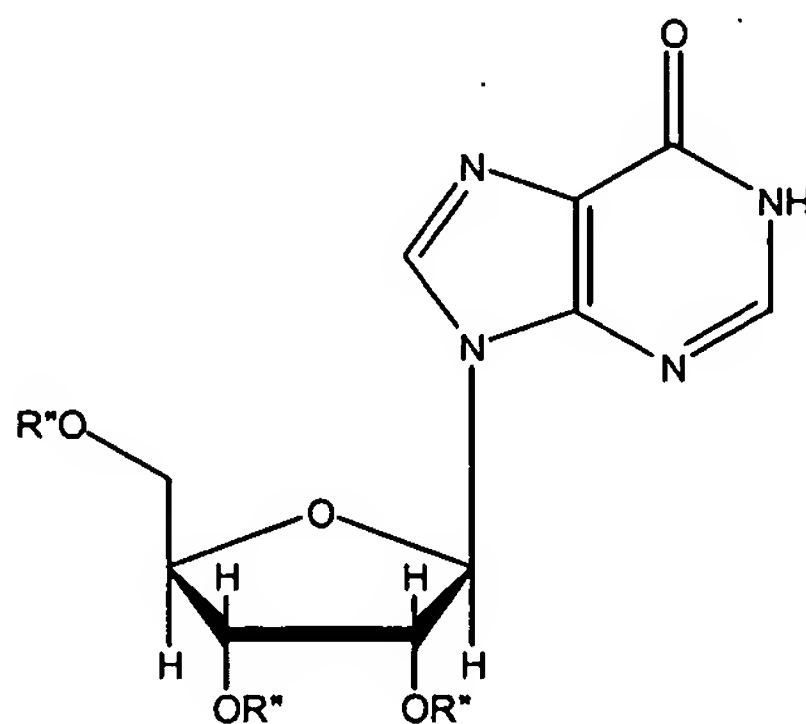
18. (Currently Amended) A method according to claim 15 ~~any of claims 15 to 17~~, wherein the compound of formula IV is triacetoxo 6-chloroadenosine, and the compound of formula III is triacetoxo 2-nitro-6-chloroadenosine.

19. (Original) A method according to claim 18, wherein triacetoxo 6-chloroadenosine is nitrated to triacetoxo 2-nitro-6-chloroadenosine using tetrabutyl ammonium nitrate (TBAN) or tetramethyl ammonium nitrate (TMAN) as nitrating reagent.

20. (Original) A method according to claim 19, which further comprises reducing the amount of tetrabutyl ammonium (TBA) or tetramethyl ammonium (TMA) impurities contaminating the triacetoxy 2-nitro-6-chloroadenosine.

21. (Original) A method according to claim 20, wherein the amount of TBA or TMA impurities is reduced by triturating the triacetoxy 2-nitro-6-chloroadenosine from isopropanol or ethanol, and washing the triturated triacetoxy 2-nitro-6-chloroadenosine with a mixture of water and ethanol.

22. (Currently Amended) A method according to claim 15 ~~any of claims 15 to 21~~, which further comprises converting a compound of formula V to a compound of formula IV:



V

wherein R'' is a protecting group, preferably acetyl or benzoyl.

23. (Original) A method according to claim 22, wherein the compound of formula V is chlorinated to form the compound of formula IV.

24. (Currently Amended) A method according to claim 22 ~~or 23~~, wherein the compound of formula V is triacetoxo inosine, and the compound of formula IV is triacetoxo 6-chloroadenosine.

25. (Original) A method according to claim 24, wherein triacetoxo inosine is chlorinated using thionyl chloride or POCl<sub>3</sub> as chlorinating reagent.

26. (Currently Amended) A method according to claim 22 ~~any of claims 22 to 25~~, which further comprises isolating the compound of formula IV produced.

27. (Currently Amended) A method according to claim 22 ~~any of claims 22 to 26~~, which further comprises converting inosine to a compound of formula V.

28. (Original) A method according to claim 27, wherein inosine is acetylated or benzoylated to form the compound of formula V.

29. (Currently Amended) A method according to claim 27 ~~or 28~~, wherein the compound of formula V is triacetoxo inosine.

30. (Original) A method according to claim 29, wherein inosine is acetylated using acetic anhydride as acetylating reagent.

31. (Currently Amended) A method according to claim 27 ~~any of claims 27 to 30~~, which further comprises isolating the compound of formula V produced.

32. (Original) A method of synthesis of spongosine which comprises the steps shown in scheme 1.

33. (Original) A method of synthesis of spongosine which is substantially as described with reference to steps 1 to 5 of the Example.

34. (Currently Amended) A 2-substituted adenosine of formula I synthesized by a method according to claim 1 ~~any of claims 1-9, or 11-33~~.

35. (Original) A method of synthesis of 2,6-dimethoxy adenosine which is substantially as described with reference to steps 1 to 4 of the Example.

36. (Currently Amended) A compound of formula II synthesized by a method according to claim 10 ~~any of claims 10 to 31, or 35~~.

37. (Original) Use of a compound of formula II, III, IV, V, triacetoxy 2-nitro, 6-chloroadenosine, triacetoxy 6-chloroadenosine, triacetoxy inosine, or inosine in the synthesis of a compound of formula I.

38. (Original) Use of a compound of formula III, IV, V, triacetoxy 2-nitro, 6-chloroadenosine, triacetoxy 6-chloroadenosine, triacetoxy inosine, or inosine in the synthesis of a compound of formula II.

39. (Original) A method of producing a nitrated substituted adenosine which comprises nitrating a substituted adenosine using TBAN or TMAN, and reducing the amount of TBA or TMA impurity contaminating the nitrated substituted adenosine.

40. (Original) A method according to claim 39, wherein the amount of TBA or TMA impurity is reduced by triturating the nitrated substituted adenosine from isopropanol or ethanol, and washing the triturated product with a mixture of water and ethanol.